

**Amendments to the Specification**

*Please amend the specification as indicated below without prejudice or disclaimer.*

*Replacement pages corresponding to these amendments are attached herewith.*

5    *Please amend lines 12-16 on page 2 as shown below:*

**Figure 1.** BFA4 cDNA sequence (SEQ ID NO.:1).

**Figure 2.** BFA4 amino acid sequence (SEQ ID NO.:2).

**Figure 3.** BCY1 nucleotide (A; SEQ ID NO.:3) and amino acid (B; SEQ ID NO.:4) sequences.

**Figure 4.** BFA5 cDNA sequence (SEQ ID NO.:5).

10    **Figure 5.** BFA5 amino acid sequence (SEQ ID NO.:6).

*Please amend the paragraph at page 14, lines 14-18 as shown below:*

A fusion motif may enhance transport of an immunogenic target to an MHC processing compartment, such as the endoplasmic reticulum. These sequences, referred to as transduction or 15 transcytosis sequences, include sequences derived from HIV tat (see Kim et al. 1997 J. Immunol. 159:1666), *Drosophila* antennapedia (see Schutze-Redelmeier et al. 1996 J. Immunol. 157:650), or human period-1 protein (hPER1; in particular, SRRHHCRSKAKRSRHH (SEQ ID NO: 105).

*Please amend Table III found on pages 30-31 as shown below:*

**TABLE III**  
***BF45 Peptide Pools***

Peptide Group	CLP number	Sequence	SEQ ID	Peptide Group	CLP number	Sequence	SEQ ID
BF45 Group 1	2983	LMDMQTFKA	7	BF45 Group 6	3033	FESSAKIQV	53
	2984	KVSIPTKAL	8		3034	GVTAEHYAV	54
	2985	SIPTKALEL	9		3035	RVTSNKTKV	55
	2986	LEIKNEQTL	10		3036	TVSQKDVCY	56
	2987	TVSQKDVCV	11		3037	KSQEPAFH	57
	2988	SVPNKAEL	12		3038	KVLAENTM	58
BF45 Group 2	2989	CETVSQKD	13	BF45 Group 7	3039	MLKLEIATL	59
	2990	KINGKLEES	14		3040	EILSWVAKL	60
	2991	SLVEKTPDE	15		3041	MILKKEIAML	61
	2992	SLCETVSQK	16		3042	LLKEKNEEI	62
	2993	EIDKINGKL	17		3043	ALRIQDIEL	63
	2994	MILQQVNVDV	18		3044	KIREELGRI	64
BF45 Group 3	2995	NMWLQQQLV	19	BF45 Group 8	3045	TLKLKEESEL	65
	2996	FLVDRKCOL	20		3046	ILNEKIREE	66
	2997	YLLHENCM	21		3047	VLKKKLSEA	67
	2998	SLFESSAKI	22		3048	GTSDDKQCL	68
	2999	KITIDIHFL	23		3049	GADINLVDV	69
	3000	QLQSKNMW	24		3050	ELCSVRLTL	70
BF45 Group 4	3001	SLDQKLFQL	25	BF45 Group 9	3051	SVESNLNQV	71
	3002	FLLIKNANA	26		3052	SLKINLNYA	72

Peptide Group	CLP number	Sequence	SEQ ID	Peptide Group	CLP number	Sequence	SEQ ID
BFA5 Group 3	3003	KILDIVHSC	<u>27</u>	BFA5 Group 8	3053	KTPDEAASL	<u>73</u>
	3004	SLSKILDIV	<u>28</u>		3054	ATCGMKVSI	<u>74</u>
	3005	ILDSSGADI	<u>29</u>		3055	LSHGVIEV	<u>75</u>
	3006	KVMEINREV	<u>30</u>		3056	EIAMLKLEI	<u>76</u>
	3007	KLLSHGAVI	<u>31</u>		3057	AELQMTLKL	<u>77</u>
	3009	AVYSEIILSV	<u>32</u>		3058	VFAADICGV	<u>78</u>
	3010	KMNVDVSVST	<u>33</u>		3060	PAIEMQNSV	<u>79</u>
	3011	ILSVVAKLL	<u>34</u>		3061	EIFNYNNHL	<u>80</u>
	3012	VLAVENTMVL	<u>35</u>		3062	ILKEKNAEL	<u>81</u>
	3013	KLSKNHQNNT	<u>36</u>	BFA5 Group 9	3063	QLVHAKKA	<u>82</u>
BFA5 Group 4	3014	SLTPLLLSI	<u>37</u>		3065	NIQDAQKRT	<u>83</u>
	3015	SQYSGQLKV	<u>38</u>		3066	NLVDVYGNM	<u>84</u>
	3016	KELEVKQQL	<u>39</u>		3067	KCTALMLAV	<u>85</u>
	3017	QIMEYIRKL	<u>40</u>		3068	KIQCLEKAT	<u>86</u>
	3018	AMLKLEIAT	<u>41</u>		3069	KIAWEKKET	<u>87</u>
	3019	VLHQPLSEA	<u>42</u>		3070	IAWEKKEDT	<u>88</u>
	3020	GLLKATCCGM	<u>43</u>		3071	VGMILLQQNV	<u>89</u>
	3021	GLLKANCAGM	<u>44</u>		3072	VKTGCVARV	<u>90</u>
	3022	QQLEQALRI	<u>45</u>	BFA5 Group 10	3074	ALHYAVYSE	<u>91</u>
	3023	CMLKKEIAM	<u>46</u>		3075	QMKKKFCVL	<u>92</u>
BFA5 Group 5	3024	EQMKKKFCV	<u>47</u>		3076	ALQCHQEAC	<u>93</u>
	3025	IQDIELKSV	<u>48</u>		3077	SEQIVEFLL	<u>94</u>
	3026	SVPNKAFFL	<u>49</u>		3078	AVIEVHNKA	<u>95</u>
	3027	SIYQKVMEI	<u>50</u>		3079	AVTCGFHHI	<u>96</u>
	3028	NLYAGDAL	<u>51</u>		3080	ACLQRKMNV	<u>97</u>
	3029	AVQDHDCIV	<u>52</u>		3081	SLVEGTSDK	<u>98</u>

*Please amend the paragraph on page 32, lines 16-32 as shown below:*

In addition to ELISPOT analysis, human T cells activated by BFA5 peptides were assayed to determine their ability to function as CTL. The cells were activated using peptide-pulsed dendritic cells followed by CD40 ligand-activated B cells (5 rounds of stimulation). The experiment shown was performed with isolated PBMC from HLA-A\*0201<sup>+</sup> donor AP31. Isolated T cells were tested in <sup>51</sup>Cr-release assays using peptide-loaded T2 cells. The % specific lysis at a 10:1, 5:1, and 1:1 T-cell to target ratio is shown for T2 cells pulsed with either pools of BFA5/NYBR-1 peptides or with individual peptides. The graph shows CTL activity induced against targets loaded with a c non-specific HLA-A\*0201-binding HIV peptide (control) followed by the CTL activity against the peptide pool (Pool 1 etc.) and then the activity induced by individual peptides from the respective pool to the right. A high level of cytotoxicity was observed for some peptides at a 1:1 E:T ratio. CTL activity (percent specific lysis) induced by the control HIV peptide was generally <10%. Similar results were obtained with another PBMC donor expressing HLA-A\*0201 (AP10). A large number of BFA5 peptides trigger T cell-mediated cytotoxicity of BFA5 peptide-loaded target cells. **Table IV** lists those peptides having immunogenic properties. Five peptides (LMDMQTFKA (SEQ ID NO.:7), ILIDSGADI (SEQ ID NO.:29), ILSVVAKLL (SEQ ID NO.:34), SQYSGQLKV (SEQ ID NO.:38), and ELCSVRLTL (SEQ ID NO.:70)) were found to induce both IFN- $\gamma$  secretion and CTL activity in T cells from both donors.

Please amend Table IV beginning on page 32, line 33 as shown below:

**TABLE IV**  
Immunoreactive peptides from BFA5

	BFA5 peptides eliciting high IFN- $\gamma$ release (>200 spots/100,000 cells)		BFA5 peptides inducing CTL lysis of pulsed cells	
<u>SEQ ID NO.</u>	Donor AP10	Donor AP31	Donor AP10	Donor AP31
<u>7</u>	LMDMQTFKA	LMDMQTFKA	LMDMQTFKA	LMDMQTFKA
<u>8</u>	KVSIPTKAL			<u>KVSIPTKAL</u>
<u>9</u>	SIPTKALEL			<u>SIPTKALEL</u>
<u>11</u>	TVSQKDVCL			
<u>12</u>	SVPNKALEL			
<u>21</u>	YLLHENCML	YLLHENCML	YLLHENCML	
<u>24</u>	QLQSKNMWL	QLQSKNMWL		QLQSKNMWL
<u>28</u>	SLSKILDV	SLSKILDV		SLSKILDV
<u>29</u>	ILIDSGADI	ILIDSGADI	ILIDSGADI	ILIDSGADI
<u>30</u>	KVMEINREV			
<u>32</u>	AVYSEILSV			
<u>34</u>	ILSVVAKLL	ILSVVAKLL	ILSVVAKLL	ILSVVAKLL
<u>37</u>	SLTPLLLSI	SLTPLLLSI		SLTPLLLSI
<u>38</u>	SQYSGQLKV	SQYSGQLKV	SQYSGQLKV	SQYSGQLKV
<u>40</u>	QIMEYIRKL	QIMEYIRKL		QIMEYIRKL
<u>49</u>	SVPNKADEL			
<u>51</u>	NLNYAGDAL	NLNYAGDAL		
<u>54</u>		GVTAEHYAV		
<u>57</u>		KSQEPAFHI		
<u>59</u>	MLKLEIATL	MLKLEIATL		MLKLEIATL
<u>61</u>		MLKKEIAML		
<u>63</u>	ALRIQDIEL			
<u>67</u>		VLKKKLSEA		
<u>70</u>	ELCSVRLTL	ELCSVRLTL	ELCSVRLTL	ELCSVRLTL
<u>72</u>	SLKINLNYA	SLKINLNYA		SLKINLNYA
<u>74</u>	ATCGMKVSI		ATCGMKVSI	
<u>77</u>	AELQMTLKL		AELQMTLKL	<u>AELQMTLKL</u>
<u>78</u>		VFAADICGV		
<u>81</u>	ILKEKNAEL	ILKEKNAEL		
<u>84</u>	NLVDVYGNM		NLVDVYGNM	
<u>85</u>	KCTALMLAV			

*Please amend lines 5-10 on page 34 as shown below:*

BFA5(1-23) KLH-MTKRKKTINLNQDAQKRTALHW (CLP-2977; SEQ ID NO:99)  
BFA5(312-334) KLH-TSEKFTWPAKGRPRKIAWEKKED (CLP-2978; SEQ ID NO:100)  
BFA5(612-634) KLH-DEILPSESKQKDYEENSWDTESL (CLP-2979; SEQ ID NO:101)  
BFA5(972-994) KLH-RLTLNQEEEKRRNADILNEKIRE (CLP-2980; SEQ ID NO: 102)  
BFA5(1117-1139) KLH-AENTMLTSKLKEKQDKEILEAEI (CLP-2981; SEQ ID NO: 103)  
BFA5(1319-1341) KLH-NYNNHLKNRIYQYEKEKAETENS (CLP-2982; SEQ ID NO: 104)

*Please amend line 26 on page 34 as shown below:*

Both bands were found to be consistent with the polyclonal antibodies tested in this analysis.